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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
10/599,451	07/18/2007	Domenico Fanara	06-796	9142	
20306 MCDONNELI	7590 05/11/200 BOEHNEN HULBER	EXAM	EXAMINER		
300 S. WACKER DRIVE			THOMAS, TIMOTHY P		
32ND FLOOR CHICAGO, IL 60606		ART UNIT	PAPER NUMBER		
		1614			
			MAIL DATE	DELIVERY MODE	
			05/11/2009	PAPER	

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Advisory Action Before the Filing of an Appeal Brief

Application No.	Applicant(s)		
10/599,451	FANARA ET AL.		
Examiner	Art Unit		
TIMOTHY P. THOMAS	1614		

y pp	Examiner	ALCOIN					
	TIMOTHY P. THOMAS	1614					
The MAILING DATE of this communication appears on the cover sheet with the correspondence address							
THE REPLY FILED 24 April 2009 FAILS TO PLACE THIS APP	PLICATION IN CONDITION FOR AL	LOWANCE.					
The reply was filed after a final rejection, but prior to or on the same day as filing a Notice of Appeal. To avoid abandonment of this application, applicant must timely file one of the following replies: (1) an amendment, affidavit, or other evidence, which places the application in condition for allowance; (2) a Notice of Appeal (with appeal fee) in compliance with 37 CFR 41.31; or (3) a Request for Continued Examination (RCE) in compliance with 37 CFR 1.114. The reply must be filed within one of the following time periods:							
The period for reply expiresmonths from the mailing The period for reply expires on: (1) the mailing date of this A no event, however, will the statutory period for reply expire is							
Examiner Note: If box 1 is checked, check either box (a) or (MONTHS OF THE FINAL REJECTION. See MPEP 706.07	Note: If box 1 is checked, check either box (a) or (b). ONLY CHECK BOX (b) WHEN THE FIRST REPLY WAS FILED WITHIN TWO OF THE FINAL REJECTION. See MPEP 706.07(f).						
Extensions of time may be obtained under 37 CFR 1.136(a). The date have been filed is the date for purposes of determining the period of ex under 37 CFR 1.17(a) is calculated from: (1) the expiration date of the set forth in (b) above, if checked. Any reply received by the Office later may reduce any earned patient term adjustment. See 37 CFR 1.704(b) NOTICE OF APPEAL	tension and the corresponding amount of shortened statutory period for reply origing than three months after the mailing date	of the fee. The appropri- nally set in the final Office	ate extension fee te action; or (2) as				
The Notice of Appeal was filed on A brief in comp	liance with 37 CFR 41 37 must be t	iled within two month	s of the date of				
filing the Notice of Appeal (37 CFR 41.37(a)), or any exter Notice of Appeal has been filed, any reply must be filed w	nsion thereof (37 CFR 41.37(e)), to	avoid dismissal of the	appeal. Since a				
<u>AMENDMENTS</u>							
 The proposed amendment(s) filed after a final rejection, I They raise new issues that would require further contains. 			cause				
 (b) They raise the issue of new matter (see NOTE below); (c) They are not deemed to place the application in better form for appeal by materially reducing or simplifying the issues for appeal; and/or 							
(d) ☐ They present additional claims without canceling a	corresponding number of finally reje	cted claims.					
NOTE: See Continuation Sheet. (See 37 CFR 1.1	16 and 41.33(a)).						
 The amendments are not in compliance with 37 CFR 1.13 		mpliant Amendment (PTOL-324).				
5. Applicant's reply has overcome the following rejection(s):							
 Newly proposed or amended claim(s) would be al non-allowable claim(s). 		•					
 For purposes of appeal, the proposed amendment(s): a) how the new or amended claims would be rejected is proving the proposed amendment of a mended claims. 		l be entered and an e	xplanation of				
The status of the claim(s) is (or will be) as follows: Claim(s) allowed:							
Claim(s) objected to: 5.							
Claim(s) rejected: <u>1.2.5.12 and 17</u> .							
Claim(s) withdrawn from consideration: 6-10,14,15 and 18 AFFIDAVIT OR OTHER EVIDENCE	<u>5-26</u> .						
The affidavit or other evidence filed after a final action, bu because applicant failed to provide a showing of good and was not earlier presented. See 37 CFR 1.116(e).	t before or on the date of filing a No d sufficient reasons why the affidavi	etice of Appeal will <u>not</u> t or other evidence is	be entered necessary and				
 The affidavit or other evidence filed after the date of filing entered because the affidavit or other evidence failed to c showing a good and sufficient reasons why it is necessary 	vercome <u>all</u> rejections under appea	l and/or appellant fail	s to provide a				
10. The affidavit or other evidence is entered. An explanatio REQUEST FOR RECONSIDERATION/OTHER	n of the status of the claims after er	ntry is below or attach	ed.				
The request for reconsideration has been considered bu See Continuation Sheet.	t does NOT place the application in	condition for allowan	ce because:				
12. Note the attached Information Disclosure Statement(s).	(PTO/SB/08) Paper No(s)						
13. Other:							
/Ardin Marschel/ Supervisory Patent Examiner, Art Unit 1614	/Timothy P Thomas/ Examiner, Art Unit 1614						
	=Adminion, Aut Offic 1014						

U.S. Patent and Trademark Office PTOL-303 (Rev. 08-06)

Continuation of 3. NOTE: the limitation of the amount of levocetirizine to less than 1 mg/mL is a new issue that requires further consideration and search.

Continuation of 11. does NOT place the application in condition for allowance because: The rejections and objections of record are maintained for the reasons of record.

Applicants argue a series of arguments based on the claim amendment, which are not relevant, since the claim amendment has not been entered.

With respect to the rejection under 35 USC 103, applicant argues that the lowest completely antibacterial concentration of the combination MP-IPP disclosed by Doron is >1.5 mg/mL, the lower concentrations tested being reported to have acterial growth, that Doron teaches that a complete antibacterial effect of a combination of parabens is not always achieved and antivacterial efficacy of parabens is weaker against planktonic bacteria compared to immobilized bacteria; therefore it is unexpected and nonoblvoit anti-ordinations according to the claims would have such antibacterial efficacy; unexpected efficacy is disclosed in Tables 15-20 of Example 4 of the present application, which have total paraben concentrations from 0.375 to 1.125 mg/mL and (MPI)[PP] = 9 are free of three bacteria types at 14 and 28 days following inoculation with these bacteria. The unexpected data is noted. However, the unexpected concentrations are not commensurate in scope with the claimed amounts, which range from greater than 0 to less than 1.5 mg/mL, in the independent claim, and only slightly narrowed in claim 5. Therefore the rejection is maintained for embodiments outside of the range for which unexpected results have been demonstrated.